GNP EXTRA STRENGTH ANTI-ITCH CREAM- diphenhydramine hydrochloride, zinc acetate cream
Amerisource Bergen

Active Ingredients

Diphenhydramine hydrochloride 2%

Zinc acetate 0.1%

Ask Doctor Section

conditions worsen

symptoms last more than 7 days or clear up and occur again within a few days

Do not use section

on chicken pox, poison ivy, sunburn, large areas of the body, broken, blistered, or oozing skin, more often than directed, or with any other product containing diphenhydramine, even one taken by mouth

If swallowed, get medical help or contact a Poison Control Center right away.

For external use only

Directions

do not use more often then directed

adults and children 2 years and older: apply to affected area not more than 3 to 4 times daily

children under 2 years: ask doctor

Uses

temporarily relieves pain and itching due to:

- insect bites
- minor burns
- sunburns
- minor skin irritations
- minor cuts
- scrapes
- rashes due to poison ivy, poison oak and poison sumac
- dries the oozing and weeping of poison:
  - Ivy
  - oak

- sumac

Store at 20C to 25C (68F to 77F)

www.mygnp.com

ceytl alcohol

diazolindinyl urea

methlparaben

polyethylene glycol

monostearate 1000

propylene glycol

propylparaben

purified water

stearyl alcohol

Diphenhydramine hydrochloride.....Topical anagesic

Zinc acetate.....Skin protectant

GNP EXTRA STRENGTH ANTI-ITCH CREAM

diphenhydramine hydrochloride, zinc acetate cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:46122-804
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	0.1 g in 100 g
DIPHENHYDRAMINE HCL (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HCL	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 1000 (UNII: U076Q6Q621)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
WATER (UNII: 059QF0KO0R)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-804-36	1 in 1 CARTON	06/01/2025	
1		1 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	06/01/2025	

Labeler - Amerisource Bergen (007914906)

Registrant - Weeks & Leo Co., Inc. (005290028)

Establishment

Name	Address	ID/FEI	Business Operations
Weeks & Leo Co., Inc		005290028	manufacture(46122-804)

Revised: 5/2025

Amerisource Bergen